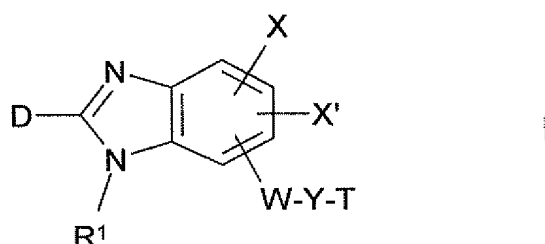


This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1-28. (Cancelled)

29. (Currently Amended) A compound of formula I ~~Compounds of the formula I according to claim 1,~~



in which

D is phenyl, pyrrolyl, furyl, thiophenyl, pyridinyl, pyrimidinyl, pyridazinyl, pyrazinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl or triazinyl, each of which is monosubstituted or polysubstituted by Hal,

X and X' are H,

W is  $-[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2CO[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nO[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nO[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2COO[C(R^2)_2]_{n-}$  or  $-[C(R^2)_2]_nS(O)_m[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,

$R^2$  is H, A or  $-[C(R^1)_2]_n-Ar'$ ,

$Ar'$  is phenyl,

Y is phenylene or piperidinediyl, each of which is unsubstituted or monosubstituted or disubstituted by A, Br, Cl or F,

T is 2-oxopiperidin-1-yl, 2-oxopyrrolidin-1-yl, 2-oxo-1*H*-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-oxo-1*H*-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxopiperazin-1-yl, 2,6-dioxopiperazin-1-yl, 2-oxopyrazin-1-yl, 2,5-dioxo-

pyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 3-oxo-2*H*-pyridazin-2-yl, 2-caprolactam-1-yl (= 2-oxoazepan-1-yl), 2-hydroxy-6-oxopiperazin-1-yl, 2-azabicyclo[2.2.2]octan-3-on-2-yl, 2-methoxy-6-oxopiperazin-1-yl, 5,6-dihydro-1*H*-pyrimidin-2-oxo-1-yl, 2-iminopiperidin-1-yl or 2-iminopyrrolidin-1-yl,

$R^1$  is H or A,

A is unbranched or branched alkyl having 1-10 carbon atoms, in which ~~and~~ 1-7 H atoms are optionally ~~may be~~ replaced by F,

m is 0, 1 or 2, and

n is 0, 1 or 2,

or a pharmaceutically acceptable derivative or solvate thereof

~~and pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios.~~

30. (Currently Amended) A compound ~~Compounds~~ according to Claim 29, which is ~~1 selected from the group consisting of~~

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyridin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyrazin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-iminopyrrolidin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-iminopiperidin-1-yl)phenyl]acetamide,

2-(5-chlorothiophen-2-yl)-5-[4-(2-oxopiperidin-1-yl)phenoxyethyl]-1*H*-benzimidazole,

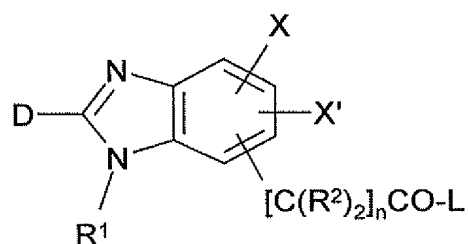
2-(5-chlorothiophen-2-yl)-5-[4-(2-oxopiperidin-1-yl)phenoxy]-1*H*-benzimidazole,  
 2-(5-chlorothiophen-2-yl)-5-[4-(2-oxopiperidin-1-yl)phenylamino]-1*H*-benzimidazole,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]valeramide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]-3-phenylpropionamide,  
 2-[2-(4-chlorophenyl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(4-chlorophenyl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chloropyridin-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chloropyridin-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]valeramide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)benzyl]acetamide,  
 1-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]formamide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-4-(2-oxopiperidin-1-yl)benzamide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylmethyl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]amine,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylamino]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-(2'-methylsulfonylbiphenyl-4-yl)acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-(3,4,5,6-tetrahydro-2*H*-

[1,4']bipyridinyl-4-ylmethyl)acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[4-(2-oxopyridin-1-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 3-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]propionamide,  
 3-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]propionamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,  
 2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-carboxamide-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]amide,  
 2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-carboxamide-*N*-[4-(3-oxomorpholin-4-yl)phenyl]amide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-iminopiperidin-1-yl)phenyl]valeramide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-oxo-2*H*-pyrazin-1-yl)phenyl]acetamide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylmethyl]-4-(2-oxopiperidin-1-yl)benzamide,  
 2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,  
 2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyridin-1-yl)phenyl]acetamide,

2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-(3,4,5,6-tetrahydro-2*H*-[1,4']bipyridinyl-4-ylmethyl)acetamide,  
2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-iminopiperidin-1-yl)phenyl]acetamide,  
2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[3-fluoro-4-(3-oxomorpholin-4-yl)phenyl]valeramide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylmethyl]-2-[4-(3-oxomorpholin-4-yl)phenyl]acetamide, or  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-2-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
or a pharmaceutically acceptable derivative or solvate thereof  
~~and pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios.~~

31. (Currently Amended) A process for preparing a compound ~~Process for the preparation of compounds of the formula I according to claim 29, comprising 1 and pharmaceutically usable derivatives, solvates and stereoisomers thereof, characterised in that~~  
a) for the preparation of a compound of the formula I in which W is  
 $-[C(R^2)_2]_nCONR^2[C(R^2)_2]_n-$ ,

reacting a compound of the formula II



## II

in which

L is Cl, Br, I or a free or reactively functionally modified OH group,  
and  $R^1$ ,  $R^2$ , D, X, X' and n are as defined for the compound of formula I in Claim 1,  
with the proviso that wherein any further OH and/or amino group present is protected,

is reacted with a compound of the formula III



III

in which

Z' is  $NHR^2[C(R^2)_2]_n$ ,

and  $R^2$ , Y, T and n are as defined for the compound of formula I in Claim 1,  
and wherein any protecting group is subsequently removed,

b) and/or converting ~~in that~~ a radical T in a compound of the formula I ~~is converted~~ into another radical T

by, for example,

- i) ~~converting a sulfanyl compound into an imino compound,~~
- ii) ~~removing an amino protecting group,~~

and/or

converting a base or acid of the compound of formula I ~~is converted~~ into one of its salts.

32. (Currently Amended) ~~Compounds of the formula I according to claim 1 as inhibitors of~~ A method for inhibiting coagulation factor Xa, comprising administering to a subject in need thereof an effective amount of a compound according to claim 29.

33. (Currently Amended) ~~Compounds of the formula I according to claim 1 as inhibitors of~~ A method for inhibiting coagulation factor VIIa, comprising administering to a subject in need thereof an effective amount of a compound according to claim 29.

34. (Currently Amended) A pharmaceutical composition, comprising a

compound according to claim 29 and a pharmaceutically acceptable carrier ~~Medicament comprising at least one compound of the formula I according to claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and optionally excipients and/or adjuvants.~~

35. (Currently Amended) A pharmaceutical composition according to claim 34, further comprising another pharmaceutically active compound other than the compound of formula I ~~Medicament comprising at least one compound of the formula I according to claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and at least one further medicament active ingredient.~~

36. (Currently Amended) Use of compounds according to claim 1 and/or physiologically acceptable salts and solvates thereof for the preparation of a medicament for the treatment of A method for treating ~~thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, a tumor, a tumor disease or tumor metastases, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 34~~ ~~tumours, tumour diseases and/or tumour metastases.~~

37 (Currently Amended) A set of kit, comprising Set (kit) consisting of separate packs of

(a) an effective amount of a compound according to claim 29, and of the formula I ~~according to claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and~~

(b) a further pharmaceutically active compound other than the compound of formula I ~~an effective amount of a further medicament active ingredient.~~

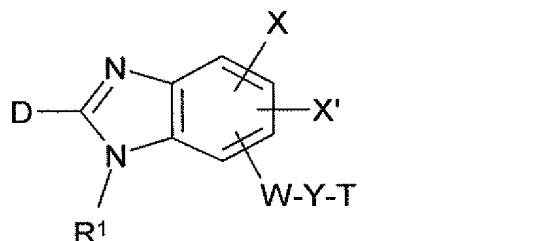
38. (Currently Amended) Use of compounds of the formula I according to claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament for the treatment of A method for treating ~~thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, a~~

tumor, a tumor disease or tumor metastases, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 35 ~~tumours, tumour diseases and/or tumour metastases, in combination with at least one further medicament active ingredient.~~

39. (New) A process according to claim 31, wherein converting a radical T in a compound of formula I into another radical T is achieved by converting a sulfanyl compound into an imino compound, or by removing an amino-protecting group.

40. (New) A compound according to claim 29, which is an isolated stereoisomer of a compound of formula I.

41. (New) A compound of formula I,



in which

D is phenyl, pyrrolyl, furyl, thiophenyl, pyridinyl, pyrimidinyl, pyridazinyl, pyrazinyl, pyrazolyl, imidazolyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl or triazinyl, each of which is monosubstituted or polysubstituted by Hal,

X and X' are H,

W is  $-[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2CO[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nO[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nO[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,  $-[C(R^2)_2]_nNR^2COO[C(R^2)_2]_{n-}$  or  $-[C(R^2)_2]_nS(O)_m[C(R^2)_2]_nCONR^2[C(R^2)_2]_{n-}$ ,

$R^2$  is H, A or  $-[C(R^1)_2]_n-Ar'$ ,

$Ar'$  is phenyl,



- Y is phenylene or piperidinediyl, each of which is unsubstituted or monosubstituted or disubstituted by A, Br, Cl or F,
- T is 2-oxopiperidin-1-yl, 2-oxopyrrolidin-1-yl, 2-oxo-1*H*-pyridin-1-yl, 3-oxomorpholin-4-yl, 4-oxo-1*H*-pyridin-1-yl, 2,6-dioxopiperidin-1-yl, 2-oxopiperazin-1-yl, 2,6-dioxopiperazin-1-yl, 2-oxopyrazin-1-yl, 2,5-dioxo-pyrrolidin-1-yl, 2-oxo-1,3-oxazolidin-3-yl, 3-oxo-2*H*-pyridazin-2-yl, 2-caprolactam-1-yl (= 2-oxoazepan-1-yl), 2-hydroxy-6-oxopiperazin-1-yl, 2-azabicyclo[2.2.2]octan-3-on-2-yl, 2-methoxy-6-oxopiperazin-1-yl, 5,6-dihydro-1*H*-pyrimidin-2-oxo-1-yl, 2-iminopiperidin-1-yl or 2-iminopyrrolidin-1-yl,
- R<sup>1</sup> is H,
- A is unbranched or branched alkyl having 1-10 carbon atoms, in which 1-7 H atoms are optionally replaced by F,
- m is 0, 1 or 2, and
- n is 0, 1 or 2,
- or a pharmaceutically acceptable salt thereof.

42. (New) A compound according to Claim 41, which is

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyridin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyrrolidin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(2-oxopyrrolidin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyrazin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-iminopyrrolidin-1-yl)phenyl]acetamide,

2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-iminopiperidin-1-

yl)phenyl]acetamide,  
 2-(5-chlorothiophen-2-yl)-5-[4-(2-oxopiperidin-1-yl)phenoxy]-1*H*-benzimidazole,  
 2-(5-chlorothiophen-2-yl)-5-[4-(2-oxopiperidin-1-yl)phenoxy]-1*H*-benzimidazole,  
 2-(5-chlorothiophen-2-yl)-5-[4-(2-oxopiperidin-1-yl)phenylamino]-1*H*-benzimidazole,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]valeramide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]-3-phenylpropionamide,  
 2-[2-(4-chlorophenyl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(4-chlorophenyl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chloropyridin-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chloropyridin-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]valeramide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)benzyl]acetamide,  
 1-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]formamide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-4-(2-oxopiperidin-1-yl)benzamide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylmethyl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]amine,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylamino]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[3-methyl-4-(3-oxomorpholin-

4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[4-(2-oxopyridin-1-yl)phenyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-sulfonyl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 3-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]propionamide,  
 3-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]propionamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,  
 2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-carboxamide-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]amide,  
 2-(5-chlorothiophen-2-yl)-1*H*-benzimidazole-5-carboxamide-*N*-[4-(3-oxomorpholin-4-yl)phenyl]amide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-iminopiperidin-1-yl)phenyl]valeramide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,  
 2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-oxo-2*H*-pyrazin-1-yl)phenyl]acetamide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylmethyl]-4-(2-oxopiperidin-1-yl)benzamide,  
 2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
 2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)phenyl]acetamide,  
 2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopyridin-1-yl)phenyl]acetamide,  
 2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[4-(2-oxopiperidin-1-yl)benzyl]acetamide,

2-[2-(5-bromothiophen-2-yl)-1*H*-benzimidazol-5-yl]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[4-(2-iminopiperidin-1-yl)phenyl]acetamide,  
2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
2-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yloxy]-*N*-[3-fluoro-4-(3-oxomorpholin-4-yl)phenyl]valeramide,  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-ylmethyl]-2-[4-(3-oxomorpholin-4-yl)phenyl]acetamide, or  
*N*-[2-(5-chlorothiophen-2-yl)-1*H*-benzimidazol-5-yl]-2-[4-(3-oxomorpholin-4-yl)phenyl]acetamide,  
or a pharmaceutically acceptable salt thereof.

43. (Currently Amended) A method for inhibiting coagulation factor Xa, comprising administering to a subject in need thereof an effective amount of a compound according to claim 41.

44. (Currently Amended) A method for inhibiting coagulation factor VIIa, comprising administering to a subject in need thereof an effective amount of a compound according to claim 41.

45. (Currently Amended) A method for inhibiting coagulation factor Xa, comprising administering to a subject in need thereof an effective amount of a compound according to claim 42.

46. (Currently Amended) A method for inhibiting coagulation factor VIIa, comprising administering to a subject in need thereof an effective amount of a compound according to claim 42.

47. (Currently Amended) A pharmaceutical composition, comprising a compound according to claim 41 and a pharmaceutically acceptable carrier

48. (Currently Amended) A pharmaceutical composition, comprising a compound according to claim 42 and a pharmaceutically acceptable carrier

49. (New) A method for treating thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, a tumor, a tumor disease or tumor metastases, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 47.

50. (New) A method for treating thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexia, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, a tumor, a tumor disease or tumor metastases, comprising administering to a subject in need thereof an effective amount of a pharmaceutical composition according to claim 48.

51. (New) A compound according to claim 41, which is an isolated stereoisomer of a compound of formula I.